INFORMATION DISCLOSURE CITATION (Use several sheets if necessary)	Attorney Docket No. 056291-5215	Application No. 10/554,202
	Applicants: **Robert Hugh BRADBURY** *et al.*	
PTO Form 1449 December 4, 2008	Filing Date: **October 24, 2005**	Group Art Unit: **1624**

U.S. PATENT DOCUMENTS

Initial		Document No.	Date	Name	Class	Sub-Class	Filing Date
	1.	US 2003/0186995	October 2, 2003	Kath et al.			
	2.	US 2004/0048880	March 11, 2004	Himmelsbach et al.			

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		Document No.	Date	Country	Class	Sub-Class	Translation
	3.	CA 2476008	October 9, 2003	Canada			
	4.	CA 2543649	May 12, 2005	Canada			
	5.	WO 01/21596	March 29, 2001	WIPO			
	6.	WO 2004/046101	June 3, 2004	WIPO			
	7.	WO 2005/041973	May 12, 2005	WIPO			
	8.	WO 2005/097134	October 20, 2005	WIPO			

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	9.	Ballard et al. "Developing a small molecule erbB2 inhibitor:challenges with optimising DMPK properties" Poster - Presented at DMDG Cambridge (February 6, 2008)
	10.	Ballard et al. "Neutral 5-substituted 4-anilinoquinazolines as potent, orally active inhibitors of erbB2 receptor tyrosine kinase" Bioorg Med Chem Lett. 17(22):6326-6329 (2007)
	11.	Barlaam et al. "A new series of neutral 5-substituted 4-anilinoquinazolines as potent, orally active inhibitors of erbB2 receptor tyrosine kinase" Bioorganic & Medicinal Chemistry Letters 18(2):674-678 (2008)
	12.	Barlaam et al. "Indazolylamino/Anilinoquinazolines Bearing a C-5 substitution as erbB2 kinase inhibitors: Structure-activity relationships and identification of a candidate drug" at AACR in 2007
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	15.	Cockerill et al. "Indazolylamino quinazolines and pyridopyrimidines as inhibitors of the EGFr and c-erbB-2" Bioorganic & Medicinal Chemistry Letters 11(11):1401-1405 (2001)
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	17.	Gaul et al. "Discovery and Biological Evaluation of Potent Dual ErbB-2/EGFR Tyrosine Kinase Inhibitors: 6-Thiazolylquinazolines" Bioorganic & Medicinal Chemistry Letters 13(4):637-640 (2003)
	18.	Harris et al. "Systematic variation of a key quinazoline core" Presented at the XXII European Colloquium on Heterocyclic Chemistry (XXII ECHC-2006) Bari, Italy, September 2-6, 2006
	19.	Hennequin et al. "N-(5-chloro-1,3-benzodioxol-4-yl)-7-[2-(4-methylpiperazin-1-yl)ethoxy]-5- (tetrahydro-2H-pyran-4-yloxy)quinazolin-4-amine, a novel, highly selective, orally available, dual-specific c-Src/Abl kinase inhibitor" J Med Chem. 49(22):6465-6488 (2006)
	20.	Jani et al. "Discovery and pharmacologic characterization of CP-724,714, a selective ErbB2 tyrosine kinase inhibitor" Cancer Research 67(20):9887-9893 (2007)

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	21.	Klutchko et al. "Tyrosine kinase inhibitors. 19. 6-Alkynamides of 4-anilinoquinazolines and 4-anilinopyrido[3,4-d]pyrimidines as irreversible inhibitors of the erbB family of tyrosine kinase receptors" J Med Chem. 49(4):1475-1485 (2006)
	22.	Petrov et al. "Optimization and SAR for dual ErbB-1/ErbB-2 tyrosine kinase inhibition in the 6-furanylquinazoline series" Bioorg Med Chem Lett. 16(17):4686-4691 (2006)

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